IN THE CLAIMS

Please cancel claims 4, 6, 14, 25-52, 58 and 60 without prejudice, and amend claims 1, 2, 5, 11, 17, 19, 21, 59 and 61, as follows:

(CURRENTLY AMENDED) A pharmaceutical composition, which is liquid, comprising:
agent i) selected from the group consisting of an insulin, an insulin analog that binds an insulin
receptor and lowers blood glucose and that differs from a naturally occurring insulin by one or more
amino acid differences, a physiologically active fragment of said insulin and a physiologically active
fragment of said insulin analog,

agent ii) selected from the group consisting of an insulin-related peptide, an insulin-related peptide analog, a physiologically active insulin-related peptide fragment and a physiologically active insulin-related peptide analog fragment, and

agent iii) an insulin sensitizer; and

a pharmaceutically acceptable non-ionic surfactant, wherein the non-ionic surfactant is a block copolymer of propylene oxide and ethylene oxide, and is present in an amount affording a concentration less than the critical micellar concentration of said composition;

wherein the insulin-related peptide is selected from the group consisting of C-peptide, glucagon-like peptide-1 (GLP-1), amylin, insulin-like growth factor-1 (IGF-1) and IGF-1 bound to binding protein 3.

- 2. (CURRENTLY AMENDED) The composition of claim 1-wherein said agent i) is further comprising an insulin.
- (ORIGINAL) The composition of claim 2 wherein said insulin is selected from the group consisting of human insulin, porcine insulin and bovine insulin.
- (CANCELLED)
- 5. (CURRENTLY AMENDED) The composition of elaim 4 claim 1 wherein said insulin analog is selected from the group consisting of Lys⁰²⁸ insulin, Pro⁰²⁹ insulin and Asp⁰²⁸ insulin.

- 6-7. (CANCELLED)
- 8. (ORIGINAL) The composition of claim 1 wherein said agent iii) is an insulin sensitizer of the glitazone family.
- 9. (ORIGINAL) The composition of claim 1 which is stabilized for administration by a medication infusion pump.
- 10. (CANCELLED)
- 11. (CURRENTLY AMENDED) The composition of claim 1, which is a liquid and comprises about 1.5 to about 40 mg/ml of agent i) and about 1.5 to about 40 mg/ml of agent ii).
- 12-16. (CANCELLED)
- 17. (CURRENTLY AMENDED) The composition of claim 1, which is a liquid and comprises about 0.5 to about 40 mg/ml of agent i) and about 0.05 to about 12 mg/ml of agent ii).
- 18. (CANCELLED)
- 19. (CURRENTLY AMENDED) The composition of claim 1, which is a liquid and comprises about 0.05 to about 12.5 mg/ml of agent ii) and about 0.05 to about 12.5 mg/ml of agent iii).
- 20. (PREVIOUSLY PRESENTED) The composition of claim 1 further comprising one or more additional compounds of agent i), of agent ii), or of agent iii).

21. (CURRENTLY AMENDED) A pharmaceutical composition comprising

- i) at least one agent selected from the group consisting of an insulin, an insulin analog, a physiologically active insulin fragment and a physiologically active insulin analog fragment and
- ii) at least one agent selected from the group consisting of an insulin-related peptide, an insulin-related peptide analog, a physiologically active insulin-related peptide fragment and a physiologically active insulated-related peptide analog fragment, and
 - iii) an insulin sensitizer; and
 - iv) optionally, a pharmaceutically acceptable carrier;

wherein said agent ii) comprises a hydrophobic portion that is coated with a pharmaceutically acceptable non-ionic surfactant that is a block copolymer of propylene oxide and ethylene oxide; wherein the composition is stabilized for administration by a medication infusion pump.

22-58. (CANCELLED)

59. (CURRENTLY AMENDED) A pharmaceutical composition comprising agents i) - iii), wherein

agent i) is selected from the group consisting of an a small molecule insulin mimetic material, agent ii) is selected from the group consisting of an insulin-related peptide, an insulin-related peptide analog, a physiologically active insulin-related peptide fragment, and a physiologically active insulin-related peptide analog fragment, and

agent iii) is an insulin sensitizer;

wherein the insulin-related peptide is selected from the group consisting of C-peptide, glucagon-like peptide-1 (GLP-1), amylin, insulin-like growth factor-1 (IGF-1) and IGF-1 bound to binding protein 3; and

wherein agents i) and ii) are combined with a pharmaceutically acceptable non-ionic surfactant that is a block copolymer of propylene oxide and ethylene oxide.

60. (CANCELLED)

61. (CURRENTLY AMENDED) The composition of claim 60 wherein the small molecule insulin mimetic material is L-783,281, having the structure:

- 62. (ORIGINAL) The composition of claim 59 wherein said agent ii) is an insulin-related peptide.
- 63. (CANCELLED)
- 64. (ORIGINAL) The composition of claim 59 wherein said agent iii) is an insulin sensitizer of the glitazone family.
- 65. (ORIGINAL) The composition of claim 59 which is stabilized for administration by a medication infusion pump.
- 66. (PREVIOUSLY PRESENTED) The composition of claim 59, which is a liquid and comprises about 1.5 to about 40 mg/ml of agent i), about 1.5 to about 40 mg/ml of agent ii), and about 0.05 to about 12.5 mg/ml of agent iii).

67-70. (CANCELLED)

71. (PREVIOUSLY PRESENTED) The composition of claim 59 further comprising one or more additional compounds of agent i), of agent ii), or of agent iii).